

DATE: Sunday, December 22, 2002 Printable Copy Create Case

Set Name side by side	Query	Hit Count Set Name result set								
DB=USPT,DWPI; PLUR=YES; OP=OR										
<u>L24</u>	L21 and maltohydrolase	0	<u>L24</u>							
<u>L23</u>	L21 and amylase	0	<u>L23</u>							
<u>L22</u>	L21 and amylase	0	<u>L22</u>							
L21	9409144	13	<u>L21</u>							
<u>L20</u>	L19 and amylase	0	<u>L20</u>							
<u>L19</u>	9205259	6	<u>L19</u>							
<u>L18</u>	9205259 and amylase	0	<u>L18</u>							
<u>L17</u>	L10 and amylase	0	<u>L17</u>							
DB=USPT; PLUR=YES; OP=OR										
<u>L16</u>	L10 and amylase	0	<u>L16</u>							
<u>L15</u>	alpha and glucan adj maltohydrolase	16	<u>L15</u>							
DB=DWPI; PLUR=YES; OP=OR										
<u>L14</u>	alpha and glucan adj maltohydrolase	0	<u>L14</u>							
<u>L13</u>	glucan adj maltohydrolase	0	<u>L13</u>							
<u>L12</u>	L10 and maltohydrolase	0	<u>L12</u>							
<u>L11</u>	L10 and beta-amylase	0	<u>L11</u>							
<u>L10</u>	9428149	3	<u>L10</u>							
DB=USPT; PLUR=YES; OP=OR										
<u>L9</u>	L8 and beta-amylase	2	<u>L9</u>							
<u>L8</u>	5034323	125	<u>L8</u>							
DB=USPT,PGPB,JPAB,EPAB,DWPI; PLUR=YES; OP=OR										
<u>L7</u>	5034323	186	<u>L7</u>							
<u>L6</u>	beta-amylase.ab. and potato and DNA	1	<u>L6</u>							
<u>L5</u>	beta-amylase.ab. and potato.ab. and DNA	0	<u>L5</u>							
<u>L4</u>	beta-amylase.ab. and potato.ab. and cDNA	0	<u>L4</u>							
<u>L3</u>	beta-amylase.ab. and potato.ab.	24	<u>L3</u>							
<u>L2</u>	beta-amylase and potato.ab.	46	<u>L2</u>							
<u>L1</u>	beta-amylase and potato	521	<u>L1</u>							

END OF SEARCH HISTORY

NEWS EXPRESS October 14 CURRENT WINDOWS VERSION IS V6.01,

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AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002

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=> s beta(w)amylase and potato 950 BETA(W) AMYLASE AND POTATO

=> s l1 and DNA

30 L1 AND DNA

=> s l1 and clone

L319 L1 AND CLONE

=> uplicate remove 13

UPLICATE IS NOT A RECOGNIZED COMMAND

The previous command name entered was not recognized by the system. For a list of commands available to you in the current file, enter

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=> n

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DUPLICATE PREFERENCE IS 'AGRICOLA, BIOSIS, EMBASE, CAPLUS'
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PROCESSING COMPLETED FOR L3

L4 9 DUPLICATE REMOVE L3 (10 DUPLICATES REMOVED)

=> d l4 1-9 ti

- L4 ANSWER 1 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE 1-
- TI Downregulation of a chloroplast-targeted ***beta*** ***amylase*** leads to a starch-excess phenotype in leaves.
- L4 ANSWER 2 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Variability in sweetpotato flavor chemistry between production years and with storage duration.
- L4 ANSWER 3 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Mass selection and root yield of true seed populations in sweet ***potato*** .
- L4 ANSWER 4 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Sites of ***beta*** ***amylase*** gene expression in sweet
 potato tuberous root: Immunohistochemistry and in situ
 hybridization studies.
- L4 ANSWER 5 OF 9 AGRICOLA DUPLICATE 2
- TI PCR cloning and sequencing of the ***beta*** ***amylase*** cDNA from barley.
- L4 ANSWER 6 OF 9 CAPLUS COPYRIGHT 2002 ACS
- TI Nucleotide sequence of a cDNA ***clone*** encoding ubiquitous .

 beta .- ***amylase*** in rye (Secale cereale L.)
- L4 ANSWER 7 OF 9 AGRICOLA DUPLICATE 3
- TI A nuclear gene encoding ***beta*** ***amylase*** of sweet ***potato*** .
- L4 ANSWER 8 OF 9 CAPLUS COPYRIGHT 2002 ACS
- TI Amylolytic activity in stored ***potato*** tubers. 2. The effect of low-temperature storage on the activities of .alpha.- and . ***beta***
 .- ***amylase*** and .alpha.-glucosidase in ***potato*** tubers
- L4 ANSWER 9 OF 9 AGRICOLA DUPLICATE 4
- TI Molecular cloning and expression in Escherichia coli of cDNA encoding the subunit of sweet ***potato*** ***beta*** ***amylase***

ANSWER 1 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE 1

ACCESSION NUMBER: DOCUMENT NUMBER:

2002:385228 BIOSIS PREV200200385228

TITLE:

Downregulation of a chloroplast-targeted ***beta***

amylase leads to a starch-excess phenotype in

leaves.

AUTHOR (S):

Scheidig, Andreas; Frohlich, Anja; Schulze, Silke; Lloyd,

James R. (1); Kossmann, Jens

CORPORATE SOURCE:

(1) Plant Biology and Biogeochemistry Department, Riso

National Laboratory, DK-4000, Roskilde: james.richard.llovd@risoe.dk Denmark

SOURCE:

Plant Journal, (June, 2002) Vol. 30, No. 5, pp. 581-591.

http://www.blackwell-science.com/

cqilib/inlpage.bin?Journal=TPJ&File=TPJ&Page=aims. print.

potato

ISSN: 0960-7412.

DOCUMENT TYPE: LANGUAGE:

Article English

AB A functional screen in Escherichia coli was established to identify ***potato*** genes coding for proteins involved in transitory starch degradation. One ***clone*** isolated had a sequence very similar to a recently described chloroplast-targeted ***beta*** - ***amylase*** of Arabidopsis. Expression of the gene in E. coli showed that the protein ***beta*** - ***amylase*** product was a functional that could degrade both starch granules and solubilized amylopectin, while import experiments demonstrated that the ***beta*** - ***amylase*** imported and processed into pea chloroplasts. To study the function of the

plants were generated where its activity was reduced using antisense

techniques. Analysis of plants reduced in the presence of this ***beta*** - ***amylase*** isoform showed that their leaves had a starch-excess phenotype, indicating a defect in starch degradation. In addition, it was shown that the antisense plants degraded only 8-30% of their total starch, in comparison with 50% in the wild type, over the dark period. This is the first time that a physiological role for a ***beta*** - ***amylase*** in plants has been demonstrated.

ANSWER 2 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. L4

protein in transitory starch degradation, transgenic

ACCESSION NUMBER: 2001:407461 BIOSIS DOCUMENT NUMBER:

PREV200100407461

TITLE:

Variability in sweetpotato flavor chemistry between

production years and with storage duration.

AUTHOR (S):

Kays, Stanley J. (1); Wang, Yan (1)

CORPORATE SOURCE:

(1) Dept. of Horticulture, The Univ. of Georgia, Athens.

GA, 30602-7273 USA

SOURCE:

Hortscience, (June, 2001) Vol. 36, No. 3, pp. 525-526.

print.

Meeting Info.: 98th Annual International Conference of the American Society for Horticultural Science Sacramento,

California, USA July 21-25, 2001

ISSN: 0018-5345.

DOCUMENT TYPE:

Conference

LANGUAGE:

English

SUMMARY LANGUAGE:

English

AB Four sweetpotato (lpomoea batatas (L.) Lam.) ***clones*** representing a diverse range in flavor types were analyzed for differences in flavor chemistry due to production year and postharvest storage duration. The

potato , Solanum (nonsweet with an aroma similar to white tuberosum L.) and 2 conventional North American cultivars, 'Jewel' and 'Centennial'. Maltose content, an indication of starch content and hydrolysis potential in baked roots, was the least variable parameter of the major sugars over 2 production years. ***clones*** Postharvest curing (7 days at 29degreeC, 95% RH) and storage duration (8 2 months at 15 degreeC, 85% RH) had a significant impact on flavor chemistry with significant quantitative variation in sugars and volatile aroma compounds in 'Jewel' and 'Centennial'. Most flavor components of GA90-156 and GA90-16 increased significantly during curing but were relatively stable during storage; neither ***clone*** produced significant maltose during baking. GA90-156 had extremely low levels of the major sugars before and during storage. Curing and storage for 2 months appeared ***beta*** - ***amylase*** , facilitating to enhance the activity of starch hydrolysis during baking and the formation of monosaccharides that act as precursors for critical volatile flavor components in 'Jewel' and 'Centennial'. Volatile aroma compounds derived from lipids, beta-carotene and terpenoids decreased with prolonged storage of 'Jewel' and 'Centennial' but did not appear to have a major qualitative impact on the flavor of the four ***clones*** . Based upon the relative low organic acid and high sugar contents of the roots, organic acids did not appear to contribute significantly to the cooked flavor of the ***clones*** tested.

L4 ANSWER 3 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 1998:362739 BIOSIS DOCUMENT NUMBER: PREV199800362739

TITLE: Mass selection and root yield of true seed populations in

sweet ***potato***

AUTHOR(S): Yoshida, Tomohiko (1)

CORPORATE SOURCE: (1) Fac. Agric., Kyushu Univ., Fukuoka 812-8581 Japan

SOURCE: Japanese Journal of Crop Science, (June, 1998) Vol. 67, No.

2, pp. 178-182. ISSN: 0011-1848.

DOCUMENT TYPE: Article LANGUAGE: Japanese

SUMMARY LANGUAGE: Japanese; English

One of the obstructions in sweet ***potato*** cultivation in temperate regions is root storage during winter for the production of vines for the next season. If a homogeneous high yielding true seed population could be developed, vine cuttings obtained from the true seed population could be transplanted in a field, thus eliminating the need for root storage. Mass selections of true seed populations were attempted to develop a high-yield and high-quality true seed population. Selection for root skin color, germination at low temperature, vine diameter and taproot diameter were effective, and a genetic gain was observed for these characteristics. Root yield obtained by transplanting vines grown from true seed population was 70 apprx 78% compared to conventionally cultured control. A ***beta*** ***amylase*** null population, which can be used for staple food and industrial purposes, was developed. The yield of true seeds by open-pollinating among ***clones*** with open field flowering was 144 seeds per ml. True seed populations with good agronomic characteristics could be used as breeding materials including overseas germplasm exchange and for an emergency crop by harvesting a large amount of true seeds, which can be stored at room temperature for a long period.

ANSWER 4 OF 9 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. L4

ACCESSION NUMBER: 1997:419979 BIOSIS DOCUMENT NUMBER: PREV199799719182

TITLE: Sites of ***beta*** - ***amvlase*** gene expression

> in sweet ***potato*** tuberous root:

Immunohistochemistry and in situ hybridization studies.

AUTHOR(S): Ling, Thai-Yen; Chou, Wen-Ling; Lee, Ping-Du; Su,

Jong-Ching

CORPORATE SOURCE: Dep. Agric. Chem., National Taiwan Univ., Taipei 10764

Taiwan

SOURCE: FASEB Journal, (1997) Vol. 11, No. 9, pp. A978.

Meeting Info.: 17th International Congress of Biochemistry

DUPLICATE 2

and Molecular Biology in conjunction with the Annual Meeting of the American Society for Biochemistry and Molecular Biology San Francisco, California, USA August

24-29, 1997 ISSN: 0892-6638.

DOCUMENT TYPE:

Conference; Abstract

LANGUAGE:

English

ANSWER 5 OF 9 AGRICOLA

ACCESSION NUMBER: 94:69210 AGRICOLA

DOCUMENT NUMBER: IND20416667

TITLE: PCR cloning and sequencing of the ***beta*** -

amylase cDNA from barley.

AUTHOR (S): Yoshigi, N.; Okada, Y.; Sahara, H.; Koshino, S.

AVAILABILITY: DNAL (385 J822)

SOURCE: The Journal of biochemistry, Jan 1994. Vol. 115, No.

1. p. 47-51

Publisher: Tokyo : Japanese Biochemical Society.

CODEN: JOBIAO; ISSN: 0021-924X

NOTE: Includes references

PUB. COUNTRY: Japan

DOCUMENT TYPE: Article

FILE SEGMENT: Non-U.S. Imprint other than FAO

LANGUAGE: English

Polymerase chain reaction (PCR) amplification of mRNA from developing barley (cultivar Haruna two-rows) endosperm was used to ***clone*** and sequence full-length cDNA encoding ***beta*** - ***amylase*** The ***beta*** - ***amylase*** cDNA was 1,775 bp in length. The ***beta*** - ***amylase*** was deduced to be composed of 535 amino acid residues and its molecular weight was calculated to be 59,610. Kreis et al. reported that the ***beta*** - ***amylase*** cDNA from barley (cultivar Hiproly) was 1,754 bp in length and coded for a polypeptide of 535 amino acids [Eur. J. Biochem. (1987) 169,517-525]. A comparison of the 3-amylase sequences from Haruna two-rows and Hiproly barleys revealed nine differences in the nucleotide sequence which resulted in three changes in the amino acid residues and 21 additional nucleotides at its 3'-end in the cultivar Haruna two-rows. The three changes were as follows: Ala-233, Ser-347, Met-527 (Haruna two-rows) and Val-233, Met-347, Ile-527 (Hiproly). Lundgard and Svensson pointed out that 23 amino acid residues of the peptide fragment derived from the COOH-terminal region of barley (cultivar Gula) ***beta*** - ***amylase*** were in agreement with the deduced amino acid sequence reported by Kreis et al., with the exception of a single position (Met-527 compared to Ile) [Carlsberg Res. Commun. (1986) 51, 487-491]. Our findings described above showed Met-527

is reasonable. In the cases of 6-amylases from soybean and sweet ***potato*** , the positions that corresponded to those at 233 and 347 in the amino acid sequence of ***beta*** - ***amylase*** from barley were Ala and Ser, respectively. Therefore, Ala-233 and Ser-347 in the amino acid sequence of barley ***beta*** - ***amylase*** were thought to be reasonable. Sequence homology of barley ***beta*** ***amylase*** with the enzymes from soybean and sweet ***potato*** amounted to 66.7 and 59.2%, respectively. ANSWER 6 OF 9 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 1994:26272 CAPLUS DOCUMENT NUMBER: 120:26272 TITLE: Nucleotide sequence of a cDNA ***clone*** ubiquitous . ***beta*** .- ***amylase*** in rye (Secale cereale L.) AUTHOR (S): Sadowski, Jan; Rorat, Tadeusz; Cooke, Richard; Delseny, Michel CORPORATE SOURCE: Inst. Plant Genet., Pol. Acad. Sci., Poznan, 60-479. Pol. SOURCE: Plant Physiology (1993), 102(1), 315-16 CODEN: PLPHAY; ISSN: 0032-0889 DOCUMENT TYPE: Journal LANGUAGE: English This report presents the sequence of a full-length cDNA encoding the ubiquitous . ***beta*** .- ***amylase*** of rye, which is the first published for monocotyledonous plants. It contains 1879 bp and a single open reading frame encoding a polypeptide of 503 amino acids. The mol. mass of the polypeptide estd. from the deduced amino acid sequence is 56,700 D. Comparison with other . ***beta*** .- ***amylase*** sequences shows 77% homol. with the partial rye endosperm-specific . ***beta*** .- ***amylase*** , 81% with the endosperm-specific form barley, 72% with that of soybean, 63% with that of Arabidopsis thaliana leaves, and 62% with that of sweet ***potato*** tuber. It has recently been shown that ubiquitous . ***beta*** .- ***amylase*** C-terminal repeat region) of wheat and rye does undergo limited proteolysis involved in conversion to the latent form during seed germination. It is probable that ubiquitous and endosperm-specific . ***beta*** .- ***amylases*** play different physiol. roles during seed development and germination. ANSWER 7 OF 9 AGRICOLA DUPLICATE 3 ACCESSION NUMBER: 93:5765 AGRICOLA DOCUMENT NUMBER: IND92075489 TITLE: A nuclear gene encoding ***beta*** - ***amylase*** of sweet ***potato*** AUTHOR (S): Yoshida, N.; Hayashi, K.; Nakamura, K. Nagoya University, Nagoya, Japan CORPORATE SOURCE: AVAILABILITY: DNAL (QH442.A1G4) SOURCE: Gene, 1992. Vol. 120, No. 2. p. 255-259 Publisher: Amsterdam : Elsevier Science Publishers. CODEN: GENED6; ISSN: 0378-1119

Includes references.

Non-U.S. Imprint other than FAO

Article

NOTE:

DOCUMENT TYPE:

FILE SEGMENT:

English

AB A nuclear AmyB gene from sweet ***potato*** encoding ***beta*** -***amylase*** (beta Amy) that is abundant in tuberous roots and inducible in other organs by an exogenous supply of sucrose or polygalacturonic acid, was isolated and characterized. Genomic Southern blot hybridization, restriction maps of independently isolated phage A ***clones*** , and the nucleotide sequence of AmyB compared with that of the cDNA, all suggested that beta Amy of sweet ***potato*** is encoded by a gene that is present in a single copy per haploid genome. In the sequence of AmyB, the sequence that is identical to that of the cDNA was split into seven exons by six introns. and the transcription of this gene starts from multiple sites 26 to 30 bp downstream from a potential TATA-box sequence, 5'-TATATAA. In the 5'-upstream region of AmyB, there are sequences homologous to those conserved in the 5'-upstream regions of genes encoding sporamin, which are regulated similarly to AmyB. The 5'-upstream region of AmyB also contains sequences to which several previously known plant nuclear factors bind.

ANSWER 8 OF 9 CAPLUS COPYRIGHT 2002 ACS T.4

ACCESSION NUMBER:

1992:425001 CAPLUS

DOCUMENT NUMBER:

117:25001

TITLE:

Amylolytic activity in stored ***potato*** tubers.

2. The effect of low-temperature storage on the activities of .alpha. - and . ***beta*** ***amylase*** and .alpha.-glucosidase in

potato tubers

AUTHOR (S):

SOURCE:

LANGUAGE:

Cochrane, M. Patricia; Duffus, Carol M.; Allison, M.

J.; Mackay, G. R.

CORPORATE SOURCE:

Edinburgh Sch. Agric., Edinburgh, EH9 3JG, UK

Potato Research (1991), 34(4), 333-41

CODEN: PORHBW; ISSN: 0014-3065

DOCUMENT TYPE:

Journal English

Tubers of the ***potato*** cultivars Record, Wilja, Pentland Dell and Brodick (formerly ***clone*** 137371) were sampled before and after storage at either 4.degree. or 10.degree.. Reducing sugar content stayed const. during storage at 10.degree. in all four cultivars but rose greatly during the first 6-12 wk of storage at 4.degree. in Record, Wilja and Pentland Dell but not in Brodick. Amylolytic activity was detd. after 5 wk storage using blocked p-nitrophenyl maltoheptaoside as substrate for .alpha.-amylase, p-nitrophenyl maltopentaoside as substrate for .

beta .- ***amylase*** , and p-nitrophenylglucopyranoside as substrate for .alpha.-glucosidase. The values obtained from tubers stored at 4.degree. were higher than those from tubers stored at 10.degree., the differences being much less in Brodick than in the other three cultivars.

ANSWER 9 OF 9 AGRICOLA

DUPLICATE 4

ACCESSION NUMBER:

92:52804 AGRICOLA

DOCUMENT NUMBER:

IND92027835

TITLE:

Molecular cloning and expression in Escherichia coli of cDNA encoding the subunit of sweet ***potato***

beta - ***amylase***

AUTHOR (S):

Yoshida, N.; Nakamaura, K.

CORPORATE SOURCE:

Mitsui Petrochemical Industries, Ltd., Waki-cho,

Kuga-gun, Yamaquchi

AVAILABILITY:

DNAL (385 J822)

SOURCE:

Journal of biochemistry, Aug 1991. Vol. 110, No. 2. p.

196-201

Publisher: Tokyo : Japanese Biochemical Society.

CODEN: JOBIAO; ISSN: 0021-924X

NOTE:

Includes references.

DOCUMENT TYPE:

Article

FILE SEGMENT:

Non-U.S. Imprint other than FAO

LANGUAGE:

English

AB Tuberous roots of the sweet ***potato*** are unusually rich in ***beta*** - ***amylase*** , and the ***beta*** - ***amylase*** polypeptides account for about 5% of the total soluble protein of the organ. Unlike ***beta*** - ***amylases*** from other origins, the identical subunits, and it also bears starch phosphorylase-inhibitor activity. A cDNA for the subunit of sweet ***potato*** ***beta*** ***amylase*** was obtained by immunological screening of an expression cDNA library constructed by the vector-primer and linker method using a plasmid vector containing tac-SP6 promoters. The SP6 transcript of a 2,000 base-pair-long cDNA insert directed the synthesis in vitro of a precursor to the subunit of ***beta*** - ***amylase*** which was identical in size with the mature subunit, and the ***beta*** - ***amylase*** mRNA detected by Northern blot hybridization was identical in size with the SP6 transcript of the cDNA insert. The cDNA insert contained 1,494 base pairs of an open reading frame which codes for the 499-amino-acid-long precursor to the subunit of ***beta*** ***amylase*** . An amino acid sequence identical to the N-terminal amino acid sequence of the mature subunit appeared immediately after the initiator methionine of the precursor, indicating that the subunit of ***beta*** - ***amylase*** is synthesized as a mature form.

Comparison

of the amino acid sequences of subunits of sweet ***potato***

beta - ***amylase*** and seed ***beta*** - ***amylases***

from barley and soybean indicated that these enzymes share about 68% amino acid identities among each other. Escherichia coli cells harboring the cDNA ***clone*** produced the mature-sized subunit of the ***beta***

- ***amylase*** , and the soluble extract exhibited amylolytic activity which migrated to the same position as the ***beta*** - ***amylase*** purified from the sweet ***potato*** in non-denaturing polyacrylamide gel containing soluble starch indicating that oligomerization of the subunit occurred properly in E. coli cells.

=> duplicate remove 12 DUPLICATE PREFERENCE IS 'AGRICOLA, BIOSIS, EMBASE, CAPLUS' KEEP DUPLICATES FROM MORE THAN ONE FILE? Y/(N):y ENTER FILE NAMES OF DUPLICATES TO KEEP:n 'N' IS NOT VALID. VALID FILE NAMES ARE 'AGRICOLA, BIOSIS, EMBASE, CAPLUS' You have entered a file name of duplicates to keep that is not referenced by any of the L#s specified for this DUPLICATE command. The file names of duplicates that can be kept are listed above. Please enter one of these file names. ENTER FILE NAMES OF DUPLICATES TO KEEP: VVV 'VVV' IS NOT VALID. VALID FILE NAMES ARE 'AGRICOLA, BIOSIS, EMBASE, CAPLUS' You have entered a file name of duplicates to keep that is not referenced by any of the L#s specified for this DUPLICATE command. The file names of duplicates that can be kept are listed above. Please enter one of these file names. ENTER FILE NAMES OF DUPLICATES TO KEEP:

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PROCESSING COMPLETED FOR L2

18 DUPLICATE REMOVE L2 AGRICOLA (12 DUPLICATES REMOVED)

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3 L5 AND TRANSFORM?

=> d 16 1-3 ibib ab

ANSWER 1 OF 3 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER:

1997:419979 BIOSIS

DOCUMENT NUMBER:

PREV199799719182

TITLE:

beta - ***amylase*** Sites of gene expression

potato tuberous root: in sweet

Immunohistochemistry and in situ hybridization studies.

AUTHOR (S):

Ling, Thai-Yen; Chou, Wen-Ling; Lee, Ping-Du; Su,

Jong-Ching

CORPORATE SOURCE:

Dep. Agric. Chem., National Taiwan Univ., Taipei 10764

Taiwan

SOURCE:

FASEB Journal, (1997) Vol. 11, No. 9, pp. A978.

Meeting Info.: 17th International Congress of Biochemistry

and Molecular Biology in conjunction with the Annual Meeting of the American Society for Biochemistry and Molecular Biology San Francisco, California, USA August

24-29, 1997 ISSN: 0892-6638.

DOCUMENT TYPE:

Conference; Abstract

LANGUAGE:

English

ANSWER 2 OF 3 CAPLUS COPYRIGHT 2002 ACS 1.6

ACCESSION NUMBER:

2000:145000 CAPLUS

DOCUMENT NUMBER:

132:190524

TITLE:

A novel plastid-targeting nucleic acid sequence, a novel . ***beta*** .- ***amylase*** sequence, and

a stimulus-responsive promoter from Arabidopsis

thaliana

INVENTOR(S):

Kavanagh, Thomas Anthony; Lao, Nga Thi

PATENT ASSIGNEE(S):

Advanced Technologies (Cambridge) Limited, UK

SOURCE:

PCT Int. Appl., 74 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIN | DATE | DATE APP | | | ON NO. | DATE | | | |
|--------------|-----------|---------|----------|-------------------------|-------|--------|----------|--------------|---|--|
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| WO 200001114 | 4 A2 | 20000 | 302 | WO 1999-GB2697 19990813 | | | | | | |
| WO 200001114 | 4 A3 | 20000 | 20000908 | | | | | | | |
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CA 2336249
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PRIORITY APPLN. INFO.:
                                        GB 1998-17959 A 19980819
                                        GB 1998-17963 A 19980819
                                        GB 1999-13014
                                                       A 19990605
                                        WO 1999-GB2697 W 19990813
     The invention provides a novel chloroplast-targeted novel . ***beta***
AΒ
     .- ***amylase*** sequence (ct . ***beta*** .- ***amylase*** ), a
     novel chloroplast targeting nucleic acid sequence, and a novel .
       ***beta*** .- ***amylase*** -encoding nucleic acid sequence from
     Arabidopsis thaliana. There is also disclosed an inducible promoter which
     is independently stimulated by light or sugar stimulus. Methods of
       ***transforming*** plants using these sequences are described, as well
          ***transformed*** plant cells, ***transformed*** plants and
     seed thereof, as well as chimeric genes contg. the sequences.
     Modification of starch levels in plants can be achieved, as well as the
     targeting of genes from the starch biosynthetic or degradative pathways,
     disease or pest resistance or variation of gene expression due to stimulus
     are described.
     ANSWER 3 OF 3 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER: 1999:753351 CAPLUS
DOCUMENT NUMBER:
                        132:946
TITLE:
                        Transgene identification in transgenic seeds using
                        screenable markers linked to aleurone-specific
                        promoters
INVENTOR(S):
                        Kriz, Alan L.; Spencer, T. Michael
PATENT ASSIGNEE(S):
                        Dekalb Genetics Corporation, USA
                        PCT Int. Appl., 177 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
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                                       WO 1999-US11023 19990518
                A1 19991125
     WO 9960129
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US 6433252 B1 20020813 US 2000-695782 20001024
PRIORITY APPLN. INFO.:
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WO 1999-US11023 W 19990518

AB The present invention provides methods and compns. for the identification of transgenic seeds. This is accomplished by use of screenable markers linked to aleurone-specific promoters. In particular embodiments of the invention, the screenable marker gene is selected from the group consisting of a green fluorescent protein gene, a luciferase gene, and an R gene; the selectable marker gene comprises a gene selected from the group consisting of neomycin phosphotransferase II, bar, EPSPS, anthranilate synthase, and dalapon dehalogenase; and the aleurone-specific promoter comprises the L3 oleosin promoter. The screenable markers can be provided as gene fusions with selectable markers, allowing both selection and screening of ***transformants*** . The use of aleurone-specific promoters, which also direct expression in embryogenic tissues, allows efficient selection of transgenic cells and the screening of viable transgenic seeds, while avoiding the deleterious effects assocd. with constitutive expression of screenable marker genes. Screening of transgenic seeds avoids the need for growing and assaying of seeds for transgenes and allows implementation of automated seed screening techniques fo the identification of transgenic seeds.

REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002

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FILE 'BIOSIS' ENTERED AT 15:11:46 ON 22 DEC 2002 COPYRIGHT (C) 2002 BIOLOGICAL ABSTRACTS INC.(R)

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L2 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2002 ACS AN 2002:408816 CAPLUS

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DN
     136:397057
ΤI
     Phosphoribosylformylglycinamidine synthase and other soybean genes
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     MacKenzie, Sally A.; Baghchhipawala, Zarir; Bassuner, Ronald
IN
PA
     The Board of Regents of the University of Nebraska, USA
SO
     PCT Int. Appl., 94 pp.
     CODEN: PIXXD2
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L_2
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AN
     2001193245 EMBASE
     Sense- and antisense-mediated gene silencing in tobacco is inhibited by
     the same viral suppressors and is associated with accumulation of small
ΑU
     Di Serio F.; Schob H.; Iglesias A.; Tarina C.; Bouldoires E.; Meins F. Jr.
     F. Meins Jr., Friedrich Miescher Institute, Novartis Research Foundation,
CS
     Maulbeerstrasse 66, CH-4058 Basel, Switzerland. meins@fmi.ch
SO
     Proceedings of the National Academy of Sciences of the United States of
     America, (22 May 2001) 98/11 (6506-6510).
     Refs: 46
     ISSN: 0027-8424 CODEN: PNASA6
CY
     United States
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FS
            Clinical Biochemistry
LA
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AN
     2001:422799 CAPLUS
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     135:208164
TI
     Copy-DNA cloning and characterization of a
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       ***glucosidase*** : expression in Escherichia coli and effects of
    down-regulation in transgenic ***potato***
ΑU
    Taylor, Mark A.; Ross, Heather A.; McRae, Diane; Wright, Frank; Viola,
    Roberto; Davies, Howard V.
CS
    Unit of Plant Biochemistry, Scottish Crop Research Institute, Dundee,
    Invergowrie, DD2 5DA, UK
SO
    Planta (2001), 213(2), 258-264
    CODEN: PLANAB; ISSN: 0032-0935
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ΤI
     A novel plastid-targeting nucleic acid sequence, a novel .beta.-amylase
     sequence, and a stimulus-responsive promoter from Arabidopsis thaliana
     Kavanagh, Thomas Anthony; Lao, Nga Thi
     Advanced Technologies (Cambridge) Limited, UK
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SO
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GB 1999-13014 A 19990605
WO 1999-GB2697 W 19990813
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AN
     1999:753351 CAPLUS
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     Transgene identification in transgenic seeds using screenable markers
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IN
     Kriz, Alan L.; Spencer, T. Michael
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     Taylor, Mark Andrew; Davies, Howard Vivian
     Nickerson Biocem Limited, UK; Taylor, Mark Andrew; Davies, Howard Vivian
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    Glucan production by ***plants*** modulation by maltooligosaccharides,
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IN
    Smith, Alison Mary; Denyer, Kay
    John Innes Centre Innovations Limited, UK; Smith, Alison Mary; Denyer, Kay
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LA English FAN.CNT 1

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     Kavanagh, Thomas Anthony; Lao, Nga Thi
IN
PΑ
     Advanced Technologies (Cambridge) Limited, UK
     PCT Int. Appl., 74 pp.
     CODEN: PIXXD2
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     132:946
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     Transgene identification in transgenic seeds using screenable markers
     linked to aleurone-specific promoters
IN
     Kriz, Alan L.; Spencer, T. Michael
PΑ
     Dekalb Genetics Corporation, USA
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     PCT Int. Appl., 177 pp.
     CODEN: PIXXD2
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                    KIND DATE
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WO 1999-US11023 19990518
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TI
     Glucan production by ***plants***
                                        modulation by maltooligosaccharides.
       ***plant*** granule-bound starch synthase I in glucan production, and
     genetic engineering to regulate maltooligosaccharide content
IN
     Smith, Alison Mary; Denyer, Kay
     John Innes Centre Innovations Limited, UK; Smith, Alison Mary; Denyer, Kay
     PCT Int. Appl., 85 pp.
     CODEN: PIXXD2
DT
     Patent
     English
FAN.CNT 1
     PATENT NO.
                  KIND DATE
                                       APPLICATION NO. DATE
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PΙ
     WO 9716554
                    A1 19970509
                                        WO 1996-GB2696 19961104
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            LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
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    AU 9673239
                     A1
                         19970522
                                       AU 1996-73239
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    EP 871744
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                                                         19961104
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    JP 11514521 T2 19991214
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L3 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2002 ACS

AB The prodn. of .alpha.-glucan (e.g. starch) by ***plant*** cells is modulated by maltooligosaccharide content in the cells. The ***plant*** granule bound starch synthase I (GBSSI) activity is regulated by maltooligosaccharides thus modulating the prodn. of unbranched .alpha.-glucan, such as amylose, and branched .alpha.-glucan, such as amylopectin. Maltooligosaccharide content of cells can be regulated by ***transforming*** ***plant*** cells with microbial .alpha.***glucosidase*** genes for example. ***Potato*** or pea
plant glucan content can thus be genetically engineered.

---Logging off of STN---

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